

https://www.who.int/health-topics/substandard-and-falsified-medical-products,

Ref. RPQ/REG/ISF/Alert N°3/2025 | | 1

All medical products must be obtained from authorized/licensed suppliers. If you have any information about the manufacture or supply of these falsified products, please contact WHO via <u>rapidalert@who.int</u>.

Annex: Products subject of WHO Medical Product Alert N°3 /2025

Product Name	IMFINZI (durvalumab) injection 500mg/10ml		
Stated manufacturer	AstraZeneca		
Identified in	Islamic Republic of Iran	Islamic Republic of Iran	Türkiye
Lot	BAZR	BBEG	AVZT
Expiry date	03-2025	12-2025	12-2026
	Availab	le Photographs	
Lot BAZR	Solut Solut Each vial Single-dos Store vials original ca Do not free	of 10 mL contains 500 mg ut durvalumab. e vial. Discard unused or non. under refrigeration at 2 0 8 8 0 in non. sze and sliake.	
Lot BBEG	Licence Holder: AstraZeneca UK Ltd., 1 Francis Crick Avenue, Cambridge Biomedical Campus, Cambridge, CB2 0AA, United Kingdom Manufactured by: Catalent Indiana, LLC, 1300 South Patterson Drive, Bloomington, IN 47403 United States of America Imported by Drimex Reg.No. EGY/BP/Aug.2019/0192/02 July all edite International Ecology Brown Market States of Edited States of America		